What is claimed:

An excipient for a metal chelate contrast agent, wherein said metal chelate contrast agent, M(L), comprises a metal ion complexed with an organic ligand, which excipient has the formula

$$X_{m}[X'(L')]_{n}$$

wherein X and X' are each independently selected from calcium or zinc, L' is an organic ligand which 10 may be L or another organic ligand which has a greater affinity for M than for calcium or zinc, and wherein m and n are each independently 1, 2 or 3.

- 15 The excipient of claim 1 wherein X = X'2. = calcium.
 - 3. The excipient of claim 1 wherein L and L' are independently selected from linear and macrocyclic polyaminopolycarboxylic acids and derivatives thereof.
 - The excipient of claim 1 wherein L and L' are independently selected from compounds of the formula

O R₂
HO-C-CH
$$CH_2$$
-CH₂
 CH_2
 CH_2

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wherein

R₁
Y is oxygen or -N-;

R₁ and R₂ are each independently hydrogen,
5 alkyl, arylalkyl, aryl, alkoxy, hydroxyalkyl,
hydroxyalkoxy,

$$(CH_2)_n$$
 NH_2 $(CH_2)_n$ R_3

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$$-(CH_{2})_{n}^{G}, (CH_{2})_{n}^{O} - C(CH_{2})_{m}^{G}, (CH_{2})_{n}^{-CH} - (CH_{2})_{m}^{G},$$

$$(CH_{2})_{n}^{G} - CH_{2}^{G} - CH_$$

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wherein G is NH_2 , NCS, $N-C-CH_2-X$, CO_2H , NHR_4 ,

 $N(R_4)_2$, CN, wherein R_4 is alkyl or hydroxyalkyl,

20 hydroxyalkoxy, -NC C $N_2^{\bigoplus}A^{\bigoplus}$ (where A is

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an anion), O-alkyl-,
$$C$$
 (CH₂)_n-SH (CH₂)_m-SH

wherein n and m are zero or an integer from one to five, R₃ is hydrogen, hydroxyalkyl, alkoxy, alkyl, aryl, arylalkyl or hydroxyalkoxy and X is chloro, bromo or iodo.

5. The excipient of claim 1 wherein L and L' are independently selected from the compounds of the formula

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$$X-CH_2$$
 CH_2-X B $V-CHR_1$ CHR_1-V

or

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 \underline{C} N(CH₂X)₃

wherein

X is -COOY, PO₃HY or -CONHOY;

Y is a hydrogen atom, a metal ion equivalent and/or a physiologically biocompatible cation of an inorganic or organic base or amino acid;

A is $-CHR_2-CHR_3-$, $-CH_2CH_2(ZCH_2-CH_2)_m-$, $N(CH_2X)_2$ $CH_2-CH_2-N(CH_2X)_2$

20 $-CH_2 - CH_2 - CH_2$, or $-CH_2 - CH_2 - CH_2$

each R₁ is hydrogen or methyl;

R₂ and R₃ together represent a trimethylene group or a tetramethylene group or individually are hydrogen atoms, lower alkyl groups (e.g., 1-8 carbons), phenyl groups, benzyl groups or R₂ is a hydrogen atom and R₃ is -(CH₂)_p-C₆H₄-W-protein where p is 0 or 1, W is -NH-, -NHCOCH₂- or -NHCS-, protein represents a protein residue;

30 m is 1, 2 or 3;

Z is an oxygen atom or a sulfur atom or the group NCH_2X or $NCH_2CH_2OR_4$ wherein X is as defined above and R_4 is C_{1-8} alkyl;

V is X or is $-CH_2OH$, $-CONH(CH_2)_nX$ or -COB, wherein X is as defined above, B is a protein or lipid residue, n is an integer from 1 to 12, or if R_1 , R_2 and R_3 are each hydrogen; then both V's together form the group

$$CH_2X$$
 CH_2X
-(CH₂)_w-N-CH₂-CH₂-N-(CH₂)_w-

- where X is as above, w is 1, 2 or 3, provided that at least two of the substituents Y represent metal ion equivalents of an element with an atomic number of 21 to 29, 42, 44 or 57 to 83.
- The excipient of claim 1 wherein L and
 L' are independently selected from the compounds of the formula

$$\underline{D} \qquad D^{5} \qquad Y-R^{2}$$
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wherein

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Y is N or P;

 ${\rm A^1}$ and ${\rm A^2}$ are each optionally branched C₂₋₆ 25 alkylene;

 ${\rm U^1}$, ${\rm U^2}$, ${\rm U^3}$ and ${\rm U^4}$ are each a single bond or optionally branched ${\rm C_{1-6}}$ alkylene;

 ${\rm D^1}$, ${\rm D^2}$, ${\rm D^3}$, ${\rm D^4}$ are each O, S, ${\rm C_{1-6}}$ alkylene or NR₇;

30 R_7 is hydrogen or C_{1-4} alkylene having a $COOR^1$ terminal group;

R1 is hydrogen or a metal ion equivalent;

 D^5 is D^1 or CHR^5 , where R^5 can be hydrogen or optionally unsaturated C_{1-20} alkylene which may include imino, phenyleneoxy, phenyleneimino, amido, ester, O, S and/or N optionally substituted with OH, SH imino and/or amino and may carry a terminal functional group (optionally bonded to a macromolecule B);

s and t are each 0-5;

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 R_2 is hydrogen, optionally substituted C_{1-16} alkyl, acyl, acylalkyl (optionally substituted by 10 one or more OH or lower alkoxy groups), $-CH_2-X-V$, B or CH_2COB where X is CO, optionally branched C_{1-10} alkylene (optionally substituted by 1 or more OH or lower alkoxy groups) or optionally branched C_{2-23} 15 alkylene interrupted by 0;

V is NR³R⁴ or COOR⁶;

 ${
m R}^3$ and ${
m R}^4$ are each hydrogen, ${
m C}_{1-16}$ alkyl (optionally substituted by 1 or more OH or lower alkoxy groups) or together complete a 5-6 membered heterocycle optionally containing another heteroatom:

R₆ is hydrogen, C₁₋₁₆ saturated, unsaturated, linear branched or cyclic hydrocarbyl, aryl or aralkyl;

 R_2 or R_3 can be bonded by a C_{2-20} alkylene 25 chain (optionally having a terminal carbonyl group, optionally interrupted by 1 or more 0 or R1 carboxymethylimino, or substituted by one or more OH, lower alkoxy or carboxy lower alkyl groups) to

30 a second macromolecule of the formula

$$\frac{D'}{D^5} \qquad \qquad \frac{D^1 - (U^2 - D^2)_S - A^1}{U^4 - D^4 - (U^3 - D^3)_+ - A^2} Y - \frac{D^4 - (U^3 - D^3)_+ - A^2}{U^4 - D^4 - (U^3 - D^3)_+ - A^2}$$

- which second macromolecule D' can be the same as or different from the macromolecule of D.
 - 7. The excipient of claim 1 wherein L and L' are independently selected from 1,4,7,10-tetra-azacyclododecane-1,4,7-triacetic acid, 1,4,7-tris-(carboxymethyl)-10-(2'-hydroxypropyl)-1,4,7,10-

tetraazacyclododecane,
N,N-bis[2-[bis(carboxymethyl)-amino]ethyl]glycine,
DTPA-bis methylamide, DTPA bis morpholinoamide and
DTPA bis 1,2-dihydroxypropylamide.

- 15 8. The excipient of claim 1 wherein L and L' are the same organic ligand.
 - 9. A contrast agent composition for use in magnetic resonance, x-ray, ultrasound and radio-diagnostic imaging comprising
- a metal ion, M, complexed with an organic ligand, L;
 - a complex salt excipient of the formula

$$X_{m}[X'(L')]_{n}$$

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wherein X and X' are each independently selected from calcium or zinc, L' is an organic ligand which may be L or another organic ligand which has a greater affinity for M than for calcium or zinc, and wherein m and n are each independently 1, 2 or 3; and,

a pharmaceutically acceptable carrier therefor.

- 10. The composition of claim 9 where \boldsymbol{X} and \boldsymbol{X}' are each calcium.
- 11. The composition of claim 9 wherein L and L' are independently selected from linear and macrocyclic polyaminopolycarboxylic acids and derivatives thereof.
- 12. The composition of claim 9 wherein L and L' are independently selected from compounds of the formula

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O R₂
HO-C-CH
$$CH_2$$
 CH_2
 CH_2

wherein

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 R_1 and R_2 are each independently hydrogen, alkyl, arylalkyl, aryl, alkoxy, hydroxyalkyl, hydroxyalkoxy,

$$(CH_2)_n \xrightarrow{C} NH_2 \xrightarrow{C} (CH_2)_n \xrightarrow{C} R_3$$

30 -
$$(CH_2)_nG$$
, $(CH_2)_n-C(CH_2)_mG$, $(CH_2)_n-CH-(CH_2)_mG$, $(CH_2)_n-CH-(CH_2)_mG$, $(CH_2)_n-CH-(CH_2)_mG$,

wherein G is NH_2 , NCS, $N-C-CH_2-X$, CO_2H , NHR_4 , H

 $N(R_4)_2$, CN, wherein R_4 is alkyl or hydroxyalkyl,

hydroxyalkoxy, -NC

N₂

N₂

N₂

(where A is

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an anion), O-alkyl- ,
$$C$$
 (CH₂)_n-SH (CH₂)_m-SH

- wherein n and m are zero or an integer from one to five, R₃ is hydrogen, hydroxyalkyl, alkoxy, alkyl, aryl, arylalkyl or hydroxyalkoxy and X is chloro, bromo or iodo.
- 13. The composition of claim 9 wherein L 20 and L' are independently selected from the compounds of the formula

$$X-CH_2$$
 CH_2-X

$$B V-CHR_1 CHR_1-V$$

or

 \underline{C} N(CH₂X)₃

wherein

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X is -COOY, PO3HY or -CONHOY;

Y is a hydrogen atom, a metal ion equivalent and/or a physiologically biocompatible cation of an inorganic or organic base or amino acid;

A is $-CHR_2-CHR_3-$, $-CH_2CH_2(ZCH_2-CH_2)_m-$, $N(CH_2X)_2$ $CH_2-CH_2-N(CH_2X)_2$ $-CH_2-CH-CH_2$, or $-CH_2-CH_2-N-CH_2-CH_2-$, wherein X is as defined above;

each R₁ is hydrogen or methyl;

R₂ and R₃ together represent a trimethylene group or a tetramethylene group or individually are hydrogen atoms, lower alkyl groups (e.g., 1-8 carbons), phenyl groups, benzyl groups or R₂ is a hydrogen atom and R₃ is -(CH₂)_p-C₆H₄-W-protein where p is 0 or 1, W is -NH-, -NHCOCH₂- or -NHCS-, protein represents a protein residue;

m is 1, 2 or 3;

Z is an oxygen atom or a sulfur atom or the group NCH_2X or $NCH_2CH_2OR_4$ wherein X is as defined above and R_4 is C_{1-8} alkyl;

V is X or is $-CH_2OH$, $-CONH(CH_2)_nX$ or -COB, wherein X is as defined above, B is a protein or lipid residue, n is an integer from 1 to 12, or if R_1 , R_2 and R_3 are each hydrogen; then both V's together form the group

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$$\begin{array}{ccc} \operatorname{CH}_2 X & \operatorname{CH}_2 X \\ -\left(\operatorname{CH}_2\right)_w - \operatorname{N-CH}_2 - \operatorname{CH}_2 - \operatorname{N-}\left(\operatorname{CH}_2\right)_w - \end{array}$$

where X is as above, w is 1, 2 or 3, provided that 30 at least two of the substituents Y represent metal ion equivalents of an element with an atomic number of 21 to 29, 42, 44 or 57 to 83. 14. The composition of claim 9 wherein L and L' are independently selected from the compounds of the formula

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$$D^{1} - D^{1} - (U^{2} - D^{2})_{S} - A^{1}$$

$$D^{5} Y - R^{2}$$

$$U^{4} - D^{4} - (U^{3} - D^{3})_{+} - A^{2}$$

wherein

10 Y is N or P;

 ${\rm A^1}$ and ${\rm A^2}$ are each optionally branched ${\rm C_{2-6}}$ alkylene;

 $\rm U^1\,,~U^2\,,~U^3$ and $\rm U^4$ are each a single bond or optionally branched $\rm C_{1-6}$ alkylene;

D¹, D², D³, D⁴ are each O, S, C₁₋₆ alkylene or NR₇;

 R_7 is hydrogen or C_{1-4} alkylene having a $COOR^1$ terminal group;

P¹ is hydrogen or a metal ion equivalent;

D⁵ is D¹ or CHR⁵, where R⁵ can be hydrogen or optionally unsaturated C₁₋₂₀ alkylene which may include imino, phenyleneoxy, phenyleneimino, amido, ester, O, S and/or N optionally substituted with OH, SH imino and/or amino and may carry a terminal functional group (optionally bonded to a macromolecule B);

s and t are each 0-5;

 R_2 is hydrogen, optionally substituted C_{1-16} alkyl, acyl, acylalkyl (optionally substituted by one or more OH or lower alkoxy groups), $-CH_2-X-V$, B or CH_2COB where X is CO, optionally branched C_{1-10} alkylene (optionally substituted by 1 or more OH or lower alkoxy groups) or optionally branched C_{2-23} alkylene interrupted by O;

V is NR3R4 or COOR6:

R³ and R⁴ are each hydrogen, C₁₋₁₆ alkyl (optionally substituted by 1 or more OH or lower alkoxy groups) or together complete a 5-6 membered heterocycle optionally containing another heteroatom:

 R_6 is hydrogen, C_{1-16} saturated, unsaturated, linear branched or cyclic hydrocarbyl, aryl or aralkyl;

10 R₂ or R₃ can be bonded by a C₂₋₂₀ alkylene chain (optionally having a terminal carbonyl group, optionally interrupted by 1 or more 0 or R¹ carboxymethylimino, or substituted by one or more OH, lower alkoxy or carboxy lower alkyl groups) to a second macromolecule of the formula

$$\frac{D'}{D^{5}} \qquad \qquad D^{5} \qquad \qquad D^{4} - D^{4} - (U^{3} - D^{3})_{+} - A^{2}$$

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which second macromolecule D' can be the same as or different from the macromolecule of D.

and L' are independently selected from 1,4,7,10-tetraazacyclododecane-1,4,7-triacetic acid, 1,4,7-tris(carboxymethyl)-10-(2'-hydroxypropyl)-1,4,7,10-tetraazacyclododecane, N,N-bis[2-[bis(carboxymethyl)-amino]ethyl]glycine, DTPA bis methylamide, 1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetic acid, DTPA bis morpholinoamide and DTPA bis 1,2-dihydroxypropylamide.

- 16. The composition of claim 9 wherein L and L' are the same organic ligand.
- 17. The composition of claim 9 wherein the mole ratio of said complex salt to said metal chelate contrast agent is between about 0.05 and 10 percent.

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- 18. The composition of claim 9 wherein said metal ion is selected from paramagnetic metal atoms, lanthanide series elements, yttrium, and the transition series elements.
- 19. The composition of claim 18 wherein said paramagnetic metals are selected from gadolinium(III), dysprosium(III), manganese(II), manganese(III), chromium(III), iron(II) and iron(III).
- 20. The composition of claim 9 wherein said metal ion complexed with an organic ligand is gadolinium(III) 1,4,7-tris(carboxymethyl)-10-(2'-hydroxypropyl)-1,4,7,10-tetraazacyclododecane and said excipient is calcium bis[1,4,7-tris(carboxymethyl)-10-(2'-hydroxypropyl)-1,4,7,10-tetraazacyclododecanatocalcium(II)].
 - 21. The composition of claim 9 wherein said metal ion complexed with an organic ligand is N-methylglucamine gadolinium (III) 1,4,7,10-tetra-azacyclododecane-N,N',N",N"'-tetraacetic acid and said excipient is calcium [1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetatocalcium(II)].
- 22. The composition of claim 9 wherein said metal ion complexed with an organic ligand is di30 N-methylglucaminium gadolinium(III) N,N-bis[2[bis(carboxymethyl)-amino]ethyl]glycine and said excipient is calcium bis[diethylenetriamineN,N',N',N",N"-pentaacetatocalcium(II)].

- 23. The composition of claim 9 wherein said metal ion complexed with an organic ligand is diethylene triamine pentaacetato-bis methylamide-gadolinium(III) and said excipient is calcium bis[diethylenetriamine-N,N'N',N",N"-pentaacetato-bis methylamide-calcium(II)].
- 24. The composition of claim 9 wherein said metal ion complexed with an organic ligand is gadolinium(III) 1,4,7,10-tetraazacyclododecane-
- 10 1,4,7-triacetic acid and said excipient is calcium bis[1,4,7,10-tetraazacyclododecane-1,4,7-triaceta-tocalcium(II)].
 - 25. The composition of claim 9 wherein said metal ion complexed with an organic ligand is gadolinium (III) DTPA bis morpholinoamide and said excipient is calcium bis [DTPA-bis morpholinamido calcium (II)].
 - 26. The composition of claim 9 wherein said metal ion complexed with an organic ligand is gadolinium (III) DTPA bis 1,2-dihydroxypropylamide and said excipient is calcium bis[DTPA bis 1,2-dihydroxypropylamido calcium (II)].
 - 27. A contrast agent composition comprising a metal chelate which is gadolinium (III)
- 1,4,7-tris(carboxymethyl)-10-(2'-hydroxypropyl)1,4,7,10-tetraazacyclododecane;

an excipient which is calcium bis[1,4,7-tris(carboxymethyl)-10-(2'-hydroxypropyl)-1,4,7,10-tetraazacyclododecanatocalcium(II)];

30 a buffer;

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acidic and/or basic solution sufficient to adjust pH of said composition to a desired value; and

water.

28. In a method of diagnostic imaging which employs an agent comprising a metal ion, M, complexed with an organic ligand, L, and a pharmaceutically acceptable carrier therefor, the improvement wherein said agent further includes an excipient of the formula

$$X_{m}[X'(L')]_{n}$$

- wherein X and X' are each independently selected from calcium or zinc, L' is an organic ligand which may be L or another organic ligand which has a greater affinity for M than for calcium or zinc, and wherein m and n are each independently 1, 2 or 3.
 - 29. The method of claim 28 wherein X and X' are each calcium.
 - 30. The method of claim 28 wherein L and L' are independently selected from linear and macrocyclic polyaminopolycarboxylic acids and derivatives thereof.
 - 31. The method of claim 28 wherein L and L' are independently selected from compounds of the formula

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O R₂
HO-C-CH
$$CH_2$$
 CH_2
 CH_2

wherein

R₁
Y is oxygen or -N-;

R₁ and R₂ are each independently hydrogen,
5 alkyl, arylalkyl, aryl, alkoxy, hydroxyalkyl,
hydroxyalkoxy,

$$(CH_2)_n$$
 NH_2 $(CH_2)_n$ R_3

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$$-(CH_2)_nG$$
, $(CH_2)_n-C(CH_2)_mG$, $(CH_2)_n-CH-(CH_2)_mG$, $(CH_2)_n-CH-(CH_2)_mG$, $(CH_2)_n-CH-(CH_2)_mG$,

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 $N(R_4)_2$, CN, wherein R_4 is alkyl or hydroxyalkyl,

hydroxyalkoxy, -N C $N_2^{\oplus}A^{\ominus}$ (where A is

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an anion), O-alkyl- ,
$$CH_2$$
)_n-SH , CH_2)_m-SH

wherein n and m are zero or an integer from one to five, R₃ is hydrogen, hydroxyalkyl, alkoxy, alkyl, aryl, arylalkyl or hydroxyalkoxy and X is chloro, bromo or iodo.

32. The method of claim 28 wherein L and L' are independently selected from the compounds of the formula

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$$X-CH_2$$
 CH_2-X B $V-CHR_1$ CHR_1-V

or

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 \underline{C} N(CH₂X)₃

wherein

X is -COOY, PO3HY or -CONHOY;

Y is a hydrogen atom, a metal ion equivalent and/or a physiologically biocompatible cation of an inorganic or organic base or amino acid;

A is $-CHR_2-CHR_3-$, $-CH_2CH_2(ZCH_2-CH_2)_m-$, $(CH_2X)_2$ $CH_2-CH_2-N(CH_2X)_2$

each R₁ is hydrogen or methyl;

R₂ and R₃ together represent a trimethylene group or a tetramethylene group or individually are hydrogen atoms, lower alkyl groups (e.g., 1-8 carbons), phenyl groups, benzyl groups or R₂ is a hydrogen atom and R₃ is -(CH₂)_p-C₆H₄-W-protein where p is 0 or 1, W is -NH-, -NHCOCH₂- or -NHCS-, protein represents a protein residue;

30 m is 1, 2 or 3;

Z is an oxygen atom or a sulfur atom or the group NCH_2X or $NCH_2CH_2OR_4$ wherein X is as defined above and R_4 is C_{1-8} alkyl;

V is X or is $-CH_2OH$, $-CONH(CH_2)_nX$ or -COB, wherein X is as defined above, B is a protein or lipid residue, n is an integer from 1 to 12, or if R_1 , R_2 and R_3 are each hydrogen; then both V's together form the group

$$\begin{array}{ccc} \operatorname{CH}_2 \operatorname{X} & \operatorname{CH}_2 \operatorname{X} \\ -\left(\operatorname{CH}_2\right)_{\mathbf{W}} - \operatorname{N} - \operatorname{CH}_2 - \operatorname{CH}_2 - \operatorname{N} - \left(\operatorname{CH}_2\right)_{\mathbf{W}} - \end{array}$$

- where X is as above, w is 1, 2 or 3, provided that at least two of the substituents Y represent metal ion equivalents of an element with an atomic number of 21 to 29, 42, 44 or 57 to 83.
- 33. The method of claim 28 wherein L and L'
 are independently selected from the compounds of
 the formula

$$\underline{D} \qquad \qquad \underline{D} \qquad \qquad \underline{D}^{5} \qquad \qquad \underline{Y} - R^{2}$$
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wherein

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Y is N or P;

 ${\tt A}^1$ and ${\tt A}^2$ are each optionally branched ${\tt C}_{2-6}$ alkylene;

 ${\rm U}^1$, ${\rm U}^2$, ${\rm U}^3$ and ${\rm U}^4$ are each a single bond or optionally branched ${\rm C}_{1-6}$ alkylene;

 $\mbox{D}^{1}\,,\mbox{ }\mbox{D}^{2}\,,\mbox{ }\mbox{D}^{3}\,,\mbox{ }\mbox{D}^{4}$ are each O, S, $\mbox{C}_{\mbox{1-6}}$ alkylene or NR $_{7}\,;$

R₇ is hydrogen or C_{1-4} alkylene having a $COOR^1$ terminal group;

R1 is hydrogen or a metal ion equivalent;

 D^5 is D^1 or CHR^5 , where R^5 can be hydrogen or optionally unsaturated C_{1-20} alkylene which may include imino, phenyleneoxy, phenyleneimino, amido, ester, O, S and/or N optionally substituted with OH, SH imino and/or amino and may carry a terminal functional group (optionally bonded to a macromolecule B);

s and t are each 0-5;

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R₂ is hydrogen, optionally substituted C₁₋₁₆
alkyl, acyl, acylalkyl (optionally substituted by
one or more OH or lower alkoxy groups), -CH₂-X-V, B
or CH₂COB where X is CO, optionally branched C₁₋₁₀
alkylene (optionally substituted by 1 or more OH or
lower alkoxy groups) or optionally branched C₂₋₂₃
alkylene interrupted by O;

V is NR3R4 or COOR6;

 ${
m R}^3$ and ${
m R}^4$ are each hydrogen, ${
m C}_{1-16}$ alkyl (optionally substituted by 1 or more OH or lower alkoxy groups) or together complete a 5-6 membered heterocycle optionally containing another heteroatom;

 R_6 is hydrogen, C_{1-16} saturated, unsaturated, linear branched or cyclic hydrocarbyl, aryl or aralkyl;

25 R₂ or R₃ can be bonded by a C₂₋₂₀ alkylene chain (optionally having a terminal carbonyl group, optionally interrupted by 1 or more 0 or R¹ carboxymethylimino, or substituted by one or more OH, lower alkoxy or carboxy lower alkyl groups) to a second macromolecule of the formula

$$\frac{D'}{D^5} = \frac{U^1 - D^1 - (U^2 - D^2)_S - A^1}{U^4 - D^4 - (U^3 - D^3)_+ - A^2}$$

- which second macromolecule D' can be the same as or different from the macromolecule of D.
- 34. The method of claim 28 wherein L and L' are independently selected from 1,4,7,10-tetra-azacyclododecane-1,4,7-triacetic acid, 1,4,7-tris-(carboxymethyl)-10-(2'-hydroxypropyl)-1,4,7,10-tetraazacyclododecane, N,N-bis[2-[bis(carboxymethyl)-amino]ethyl]glycine, DTPA-bis methylamide, 1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetic acid, DTPA bis morpholinoamide and DTPA bis 1,2-dihyroxypropylamide.
 - 35. The method of claim 28 wherein L and L' are the same organic ligand.
- 36. The method of claim 28 wherein the mole ratio of said complex salt to said metal chelate contrast agent is between about 0.05 and 10 percent.

- 37. The method of claim 28 wherein said metal ion is selected from paramagnetic metal atoms, lanthanide series elements, yttrium, and the transition series elements.
- 38. The method of claim 28 wherein said paramagnetic metals are selected from gadolinium (III), octahedral manganese(II), chromium(III), and iron(III).

39. The method of claim 28 wherein said metal ion complexed with an organic ligand is gadolinium(III) 1,4,7-tris(carboxymethyl)-10-(2'-hydroxypropyl)-1,4,7,10-tetraazacyclododecane and said excipient is calcium bis[1,4,7-tris(carboxymethyl)-10-(2'-hydroxypropyl)-1,4,7,10-tetraazacyclododecanatocalcium(II)].

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- 40. The method of claim 28 wherein said metal ion complexed with an organic ligand is N-methylglucamine gadolinium (III) 1,4,7,10-tetra-azacyclododecane-N,N',N",N"'-tetraacetic acid and said excipient is calcium [1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetatocalcium(II)].
- 41. The method of claim 28 wherein said

 metal ion complexed with an organic ligand is diNmethylglucamine gadolinium(III) N,N-bis[2-[bis(carboxymethyl)-amino]ethyl]glycine and said
 excipient is calcium bis[diethylenetriamineN,N',N',N",N"-pentaacetatocalcium(II)].
- 42. The method of claim 28 wherein said metal ion complexed with an organic ligand is gadolinium(III) N,N-bis[2-[bis(carboxymethyl)-amino]ethyl]glycine-bis methylamide and said excipient is calcium bis[diethylenetriamine-N,N'N',N",N"-pentaacetato-bis methylamide-calcium(II)].

- 43. The method of claim 28 wherein said metal ion complexed with an organic ligand is gadolinium(III) 1,4,7,10-tetraazacyclododecane-1,4,7-triacetic acid and said excipient is calcium bis[1,4,7,10-tetraazacyclododecane-1,4,7-triaceta-tocalcium(II)].
- 44. The method of claim 28 wherein said metal ion complexed with an organic ligand is gadolinium (III) DTPA bis morpholinoamide and said excipient is calcium bis [DTPA-bis morpholinamido calcium (II)].
- 45. The method of claim 28 wherein said metal ion complexed with an organic ligand is gadolinium (III) DTPA bis 1,2-dihydroxypropylamide and said excipient is calcium bis[DTPA bis 1,2-dihydroxypropylamido calcium (II)].
- 46. The method of claim 28 wherein said contrast agent composition comprising
- a metal chelate which is gadolinium (III)

 1,4,7-tris(carboxymethyl)-10-(2'-hydroxypropyl)
 1,4,7,10-tetraazacyclododecane;

an excipient which is calcium bis[1,4,7-tris(carboxymethyl)-10-(2'-hydroxypropyl)-1,4,7,10-tetraazacyclododecanatocalcium(II)];

a buffer;

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acidic and/or basic solution sufficient to adjust pH of said composition to a desired value; and

water.